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                 with preparation role
NEWS
         DEC 18
      4
                 CA/CAplus patent kind codes updated
NEWS
     5
         DEC 18
                 MARPAT to CA/CAplus accession number crossover limit increased
                 to 50,000
                 MEDLINE updated in preparation for 2007 reload
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         DEC 18
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     7
         DEC 27
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         JAN 08
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         JAN 16
                 CA/CAplus Company Name Thesaurus enhanced and reloaded
NEWS 10
         JAN 16
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         JAN 16
                 WPIDS/WPINDEX/WPIX enhanced with IPC 8 reclassification data
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         JAN 22
                 CA/CAplus updated with revised CAS roles
NEWS 13
         JAN 22
                 CA/CAplus enhanced with patent applications from India
NEWS 14
         JAN 29
                 PHAR reloaded with new search and display fields
NEWS 15
         JAN 29
                 CAS Registry Number crossover limit increased to 300,000 in
                 multiple databases
NEWS 16
         FEB 15
                 PATDPASPC enhanced with Drug Approval numbers
         FEB 15
NEWS 17
                 RUSSIAPAT enhanced with pre-1994 records
                 KOREAPAT enhanced with IPC 8 features and functionality
         FEB 23
NEWS 18
NEWS 19
         FEB 26
                 MEDLINE reloaded with enhancements
                 EMBASE enhanced with Clinical Trial Number field
NEWS 20
         FEB 26
NEWS 21
         FEB 26
                 TOXCENTER enhanced with reloaded MEDLINE
NEWS 22
         FEB 26
                 IFICDB/IFIPAT/IFIUDB reloaded with enhancements
NEWS 23
         FEB 26
                 CAS Registry Number crossover limit increased from 10,000
                 to 300,000 in multiple databases
NEWS 24
         MAR 15
                 WPIDS/WPIX enhanced with new FRAGHITSTR display format
NEWS 25
         MAR 16
                 CASREACT coverage extended
NEWS 26
         MAR 20
                 MARPAT now updated daily
NEWS 27
         MAR 22
                 LWPI reloaded
NEWS 28
         MAR 30
                 RDISCLOSURE reloaded with enhancements
NEWS 29
         MAR 30
                 INPADOCDB will replace INPADOC on STN
NEWS 30
         APR 02
                 JICST-EPLUS removed from database clusters and STN
NEWS EXPRESS NOVEMBER 10 CURRENT WINDOWS VERSION IS V8.01c, CURRENT
              MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP),
              AND CURRENT DISCOVER FILE IS DATED 25 SEPTEMBER 2006.
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=> file reg
COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE TOTAL ENTRY SESSION 0.21 . 0.21

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=>
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L1 STRUCTURE UPLOADED

=> d l1

L1 HAS NO ANSWERS

L1 STR

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

Structure attributes must be viewed using STN Express query preparation.

=> s 11

Erich Leeser

SAMPLE SEARCH INITIATED 16:11:23 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 441216 TO ITERATE

0.5% PROCESSED 2000 ITERATIONS 5 ANSWERS

INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**

BATCH **INCOMPLETE**

PROJECTED ITERATIONS: 8786458 TO 8862182
PROJECTED ANSWERS: 20068 TO 24052

L2 5 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 16:11:27 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 8819948 TO ITERATE

4.6% PROCESSED 408381 ITERATIONS 1453 ANSWERS

9.5% PROCESSED 839026 ITERATIONS 4501 ANSWERS

4878 ANSWERS

11.3% PROCESSED 1000000 ITERATIONS

INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)

SEARCH TIME: 00.00.46

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**

BATCH **INCOMPLETE**

PROJECTED ITERATIONS: 8819948 TO 8819948 PROJECTED ANSWERS: 42401 TO 43645

L3 4878 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS SINCE FILE TOTAL ENTRY SESSION

FULL ESTIMATED COST 173.90 174.11

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http://www.cas.org/infopolicy.html

=> s 13

L4 158 L3

=> s 13 full

L5 158 L3

=> s 15 and py<2003

22870367 PY<2003

L6 2 L5 AND PY<2003

=> d ibib abs hitstr tot

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 2006:740618 CAPLUS

DOCUMENT NUMBER: 145:159828

TITLE: Nicotine in therapeutic angiogenesis and

vasculogenesis

INVENTOR(S): Cooke, John; Jang, James; Tsao, Phillip; Heeschen,

Christopher

PATENT ASSIGNEE(S): USA

SOURCE: U.S. Pat. Appl. Publ., 22 pp., Cont.-in-part of U.S.

Ser. No. 147,389.

CODEN: USXXCO

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 4

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
	,			
US 2006167028	A1	20060727	US 2005-286850	20051122
US 6417205	В1	20020709	US 2000-628226	20000728 <
US 2002128294	A1	20020912	US 2002-147389	20020515 <
US 7160904	B2	20070109		
US 2006182731	A1	20060817	US 2006-404445	20060413
PRIORITY APPLN. INFO.:			US 1999-146233P	P 19990728
			US 2000-628226	A3 20000728
			US 2002-147389	A2 20020515
			US 2005-286850	A1 20051122

AB The invention features methods for induction of angiogenesis by administration of nicotine or other nicotine receptor agonist. Induction of angiogenesis by the methods of the invention can be used in therapeutic angiogenesis in, for example, treatment of ischemic syndromes such as coronary or peripheral arterial disease. Nicotine stimulated angiogenesis in mice implanted with the disk angiogenesis system and this effect was blocked by nitric oxide and prostacyclin inhibitors. Nicotine enhanced angiogenesis in a murine model of peripheral artery disease.

IT 900492-78-6

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(nicotine for therapeutic angiogenesis and vasculogenesis in ischemic syndromes)

RN 900492-78-6 CAPLUS

CN 1H-Cyclopenta[b]quinolin-1-one, 2,3-dihydro-9-methoxy-2-[2-[1-(phenylmethyl)-4-piperidinyl]ethyl]- (9CI) (CA INDEX NAME)

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2007 ACS on STN

ACCESSION NUMBER: 1966:35721 CAPLUS

DOCUMENT NUMBER: 64:35721

ORIGINAL REFERENCE NO.: 64:6599q-h,6600a-b

TITLE: Enols in the pyrrolidine series. Reaction site and

stereoselectivity in reactions of certain aluminum and

boron hydrides with α, β -unsaturated ketones

and their enolic reduction products

AUTHOR(S): Southwick, Philip L.; Latif, Nazih; Fitzgerald,

Berenice M.; Zaczek, Norbert M.

CORPORATE SOURCE: Carnegie Inst. of Technol., Pittsburgh, PA

SOURCE: Journal of Organic Chemistry (1966), 31(1),

1-1

CODEN: JOCEAH; ISSN: 0022-3263

DOCUMENT TYPE: Journal LANGUAGE: English

OTHER SOURCE(S): CASREACT 64:35721
GI For diagram(s), see printed CA Issue.

AB A study of the action of several hydride reducing agents and Grignard reagents on the α,β -unsatd. ketonic structure of

4-benzylidene-2,3-dioxopyrrolidines (I) led to the following conclusions. (1) The. initial reaction of diborane involves the olefinic bond and leads to formation of enols (II) of 4-benzyl-2,3-dioxopyrrolidines, as does catalytic hydrogenation. A comparison with chalcone showed that the latter substance is attacked to a considerable extent at the olefinic bond (conjugated reduction to the saturated ketone) by lithium tri-tertbutoxyaluminohydride (LBAH) as well as by diborane. (2) The initial reaction of LBAH, sodium borohydrde, PhMgBr, MeMgI, and presumably of LiAlH4 with I takes place at the ketonic carbonyl and produces the 4-benzylidene-3-hydroxypyrrolidine derivs. III, IV, or V. (3) III are isomerized rapidly at room temperature to II by sodium hydroxide, LBAH, or sodium borohydride. At 0° the isomerization is too slow to interfere seriously with isolation of III formed in LBAH or sodium borohydride redns. (4) II are reduced by LiAlH4 and sodium borohydride but not by LBAH or diborane. (5) Stereoselectivity in hydride redns. of II favors products having the 4-benzyl trans to the 3-hydroxyl. The same stereoisomers are favored in redns. of V with LiAlH4 at high temps.

IT 905825-22-1P, 2-Pyrrolidinone, 1-cyclohexyl-4-[p-

(dimethylamino)benzyl]-3-hydroxy-, trans-

RL: PREP (Preparation)

(preparation of)

RN 905825-22-1 CAPLUS

Relative stereochemistry.